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(54) Photographic silver halide materials and process comprising new pyrazoloazole coupler.

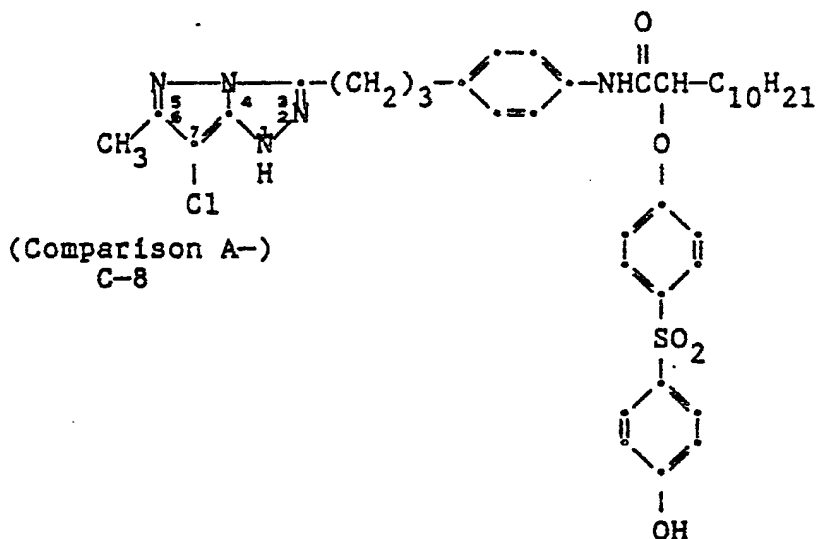
(57) Novel pyrazoloazole dye-forming couplers contain on theazole ring a group containing a carbon atom (A) bonded directly to theazole ring and wherein this carbon atom (A) is bonded directly to a nitrogen atom in the group. Such couplers provide advantageous bathochromic hue shift in dyes formed from the couplers. These couplers are useful in photographic silver halide materials and processes.

EP 0 285 274 A1

# PHOTOGRAPHIC SILVER HALIDE MATERIALS AND PROCESS COMPRISING NEW PYRAZOLOAZOLE COUPLER

This invention relates to novel pyrazoloazoles as dye-forming couplers and to photographic silver halide elements and processes using such couplers.

Color images are customarily obtained in the photographic art by reaction between the oxidation product of a silver halide color developing agent and a dye-forming coupler. Pyrazolone couplers are useful for forming magenta dye images, however, pyrazoloazole couplers represent another class of couplers that are useful for this purpose. Examples of pyrazolotriazole couplers, particularly 1H-pyrazolo [3,2-c]-s-triazole couplers, are described in, for example, U.S Patent 4,443,536; U.K. Patents 1,247,493; 1,252,418 and 1,398,979. An example of such a coupler is represented by the formula:

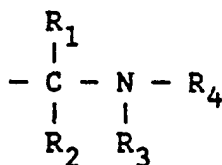


While such couplers are useful in photographic silver halide materials and processes, many such couplers have not had the desired degree of dispersibility and photographic activity to provide increased maximum dye density. Additionally, they do not always provide dyes of the desired hue.

It has been desirable to provide pyrazoloazole couplers that enable the desired bathochromic hue shift without undesired reduction in photographic activity and dispersibility.

The present invention provides a photographic silver halide element comprising a support bearing at least one photographic silver halide emulsion layer and in or adjacent the emulsion layer a pyrazoloazole color coupler having substituted on the azole ring of the coupler a group containing a carbon atom (A) bonded directly to the azole ring and wherein the carbon atom (A) is bonded directly to a nitrogen atom in the group that enables a bathochromic hue shift in a dye formed from the pyrazoloazole coupler.

The pyrazoloazole coupler preferably has substituted on the azole ring of the coupler a group represented by the structure:

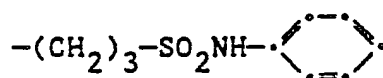
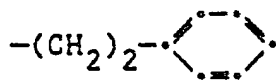
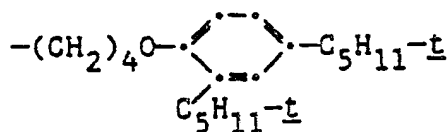


wherein

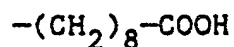
$R_1$  and  $R_2$  are individually hydrogen, unsubstituted alkyl, such as alkyl containing 1 to 40 carbon atoms, for example methyl, ethyl, propyl, butyl, pentyl and eicosyl; substituted alkyl, such as substituted alkyl containing 1 to 40 carbon atoms, for example alkyl substituted with a water solubilizing group, such as carboxy or hydroxy; unsubstituted aryl, such as aryl containing 6 to 40 carbon atoms, for example phenyl or

naphthyl, or substituted aryl, such as substituted aryl containing 7 to 40 carbon atoms, for example, methoxyphenyl or xylyl; and other substituents that do not adversely affect the coupler;

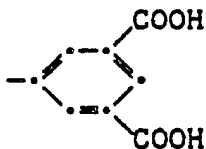
$R_1$  and  $R_2$  are individually hydrogen, unsubstituted alkyl, such as alkyl containing 1 to 40 carbon atoms, for example, methyl, ethyl, propyl, butyl, pentyl, decyl, and eicosyl; substituted alkyl, such as substituted alkyl containing 1 to 40 carbon atoms, for example,

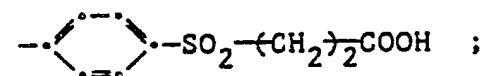
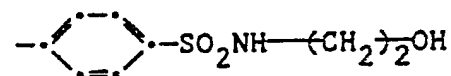
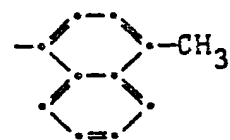
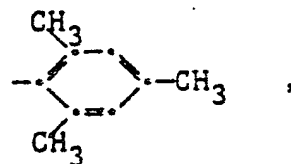
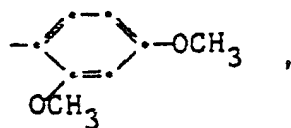


or



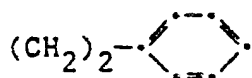
unsubstituted aryl, such as aryl containing 6 to 40 carbon atoms, for example, phenyl and naphthyl; or substituted aryl, such as substituted aryl containing 7 to 40 carbon atoms, for example, carboxyphenyl,





or  $\text{C}(=\text{O})\text{---R}_3$ ; wherein at least one of  $\text{R}_3$  and  $\text{R}_4$  is unsubstituted or substituted alkyl,  
aryl or  $\text{C}(=\text{O})\text{---R}_5$ ;

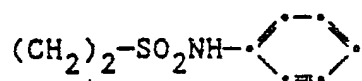
wherein  $\text{R}_5$  is alkyl, such as alkyl containing 1 to 40 carbon atoms, for example, unsubstituted alkyl, such as methyl, ethyl, propyl, butyl, octyl, and eicosyl or substituted alkyl, such as substituted alkyl containing 2 to 40 carbon atoms, for example,



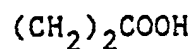
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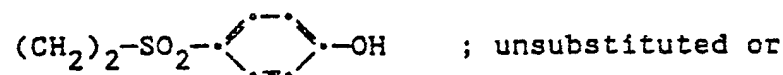
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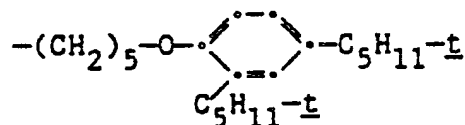


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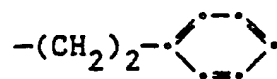
or  $\text{NHR}_s$ ,

wherein  $R_s$  is a substituent that does not adversely affect the coupler, such as alkyl, for example, alkyl containing 1 to 40 carbon atoms, such as unsubstituted or substituted alkyl, for example, methyl, ethyl, propyl, butyl, octadecyl,

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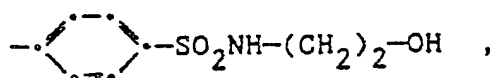
$-(\text{CH}_2)_2\text{OH}$  and  $-(\text{CH}_2)_2\text{COOH}$  ;

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unsubstituted or substituted aryl, such as aryl containing 6 to 40 carbon atoms, for example, phenyl, naphthyl, carboxyphenyl, hydroxyphenyl,

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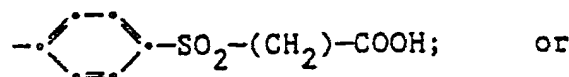
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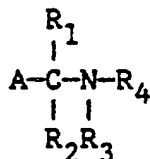
<sup>15</sup> -COOR<sub>7</sub>, wherein R<sub>7</sub> is unsubstituted or substituted alkyl, such as ethyl, propyl, octyl, or eicosyl; or unsubstituted or substituted aryl, such as phenyl, naphthyl, carboxyphenyl or xylyl. The carbon atom in the group containing R<sub>1</sub> and R<sub>2</sub> is carbon atom (A). The above group containing R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub> and R<sub>4</sub> preferably comprises a ballast group. The pyrazoloazole coupler preferably comprises at least one water solubilizing group, such as a carboxy group.

20 These pyrazolazole couplers not only provide a desired bathochromic shift in hue and enable increased maximum dye density but also enable formation of improved dispersions without adverse effects on desired properties, such as sensitivity of the photographic material.

The pyrazoloazole couplers are typically pyrazolotriazoles or pyrazolodiazoles; however, other pyrazoloazole couplers are also useful. Combinations of pyrazoloazole couplers according to the invention are also useful.

Particularly useful pyrazolotriazoles are pyrazolo [3,2-c]-s-triazoles and pyrazolo [2,3-b]-s-triazoles. The pyrazolo [2,3-b]-s-triazole can also be named as a pyrazolo [1,5-b]-1,2,4-triazole.

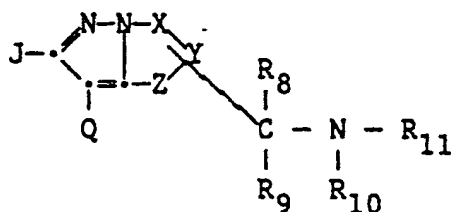
A typical pyrazoloazole coupler according to the invention is represented by the formula:



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wherein A is a pyrazoloazole coupler nucleus and, R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub>, and R<sub>4</sub> are as described. R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub> and/or R<sub>4</sub> preferably contain at least one water solubilizing group, such as a carboxy group.

A typical pyrazolotriazole coupler according to the invention is represented by the structure:



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50 wherein:

J is a substituent which does not adversely affect the desired properties of the coupler; particularly unsubstituted or substituted alkyl or unsubstituted or substituted aryl;

Q is a hydrogen or a coupling-off group such as halogen or a coupling-off group known to be useful for 2-equivalent couplers in the photographic art;

55 X,Y and Z are individually selected from carbon and nitrogen atoms necessary to complete an azole ring, particularly a triazole ring with unsaturated bonding being present in the ring as needed;

R<sub>4</sub> and R<sub>5</sub> are individually hydrogen or unsubstituted or substituted alkyl; or unsubstituted or substituted aryl, as defined for R<sub>1</sub> and R<sub>2</sub>;

R<sub>10</sub> and R<sub>11</sub> are individually hydrogen, unsubstituted or substituted alkyl; unsubstituted or substituted aryl or

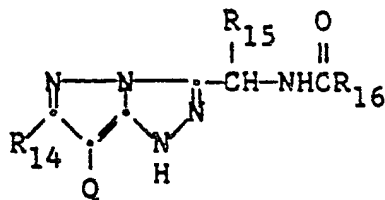
$\text{C}(=\text{O})-\text{R}_{12}$ , as defined for R<sub>1</sub> and R<sub>4</sub>; wherein at least one

of R<sub>10</sub> and R<sub>11</sub> is alkyl, aryl or  $\text{C}(=\text{O})-\text{R}_{12}$ ;

R<sub>12</sub> is unsubstituted or substituted alkyl, unsubstituted or substituted aryl or NH-R<sub>13</sub>; and

R<sub>13</sub> is unsubstituted or substituted alkyl or unsubstituted or substituted aryl.

Preferred 1H-pyrazolo[3,2-c]-s-triazole couplers are represented by the formula



wherein:

R<sub>14</sub> is a substituent that does not adversely affect the desired properties of the pyrazolotriazole coupler particularly unsubstituted or substituted alkyl or unsubstituted or substituted aryl;

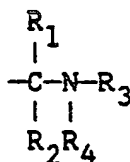
R<sub>15</sub> is unsubstituted or substituted alkyl;

R<sub>16</sub> is unsubstituted or substituted alkyl or NH-R<sub>17</sub>;

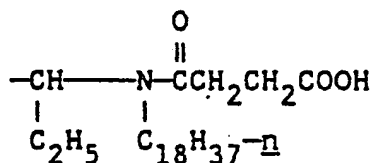
R<sub>17</sub> is unsubstituted or substituted alkyl, or unsubstituted or substituted aryl; and

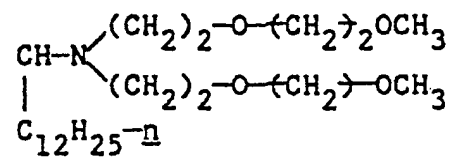
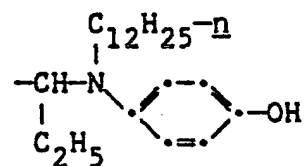
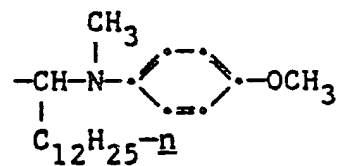
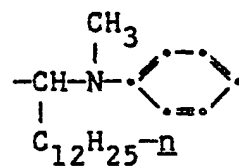
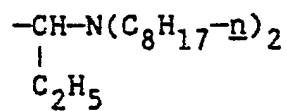
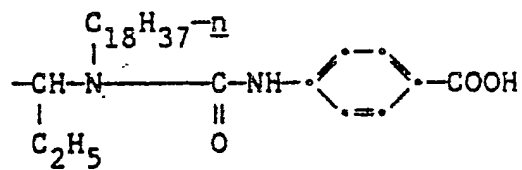
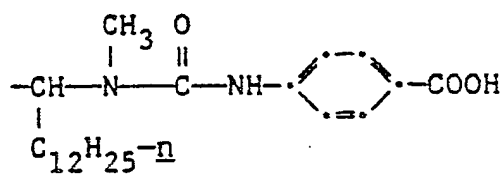
Q is hydrogen or a coupling-off group.

Illustrative examples of the group



are:

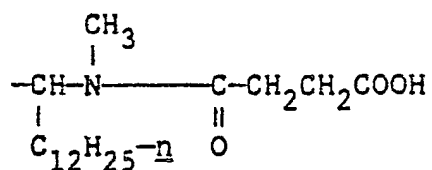




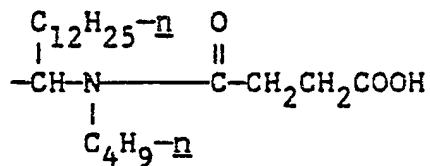




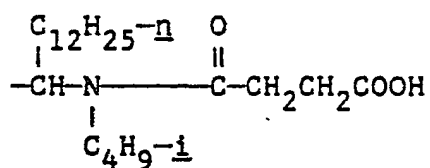
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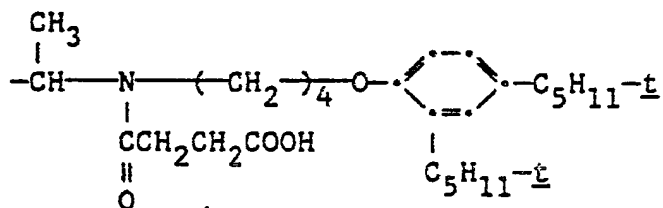


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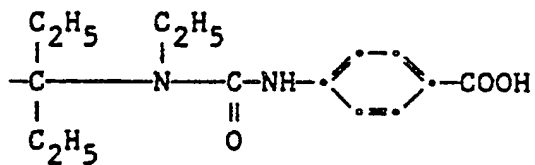
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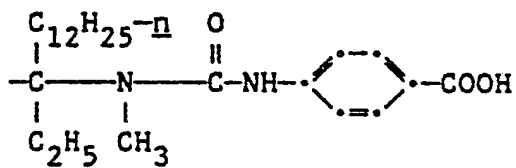


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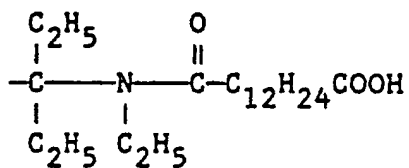
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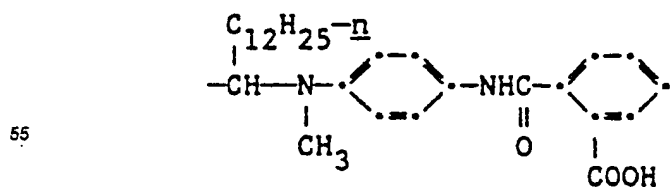
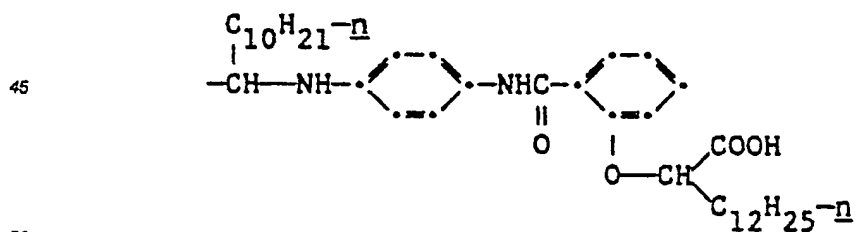
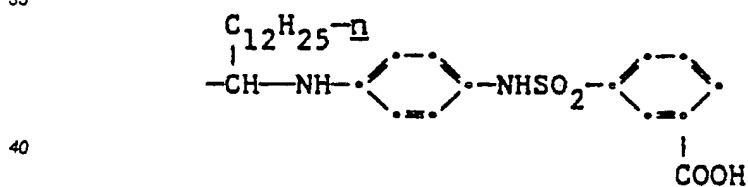
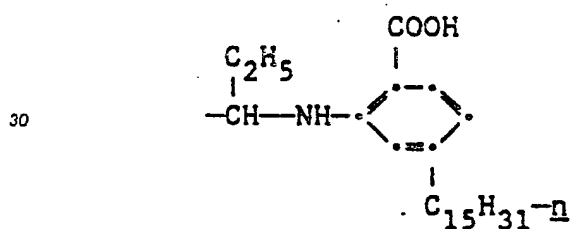
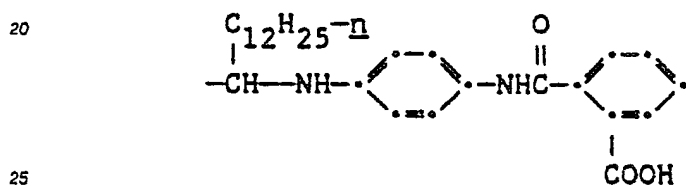
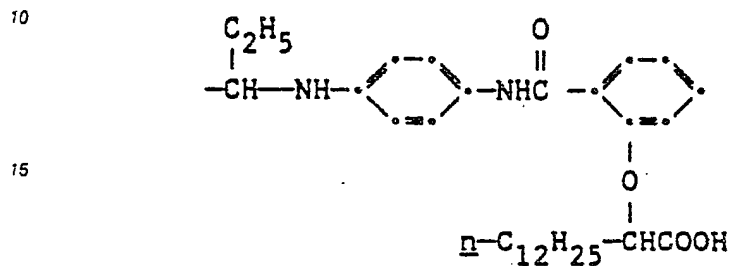
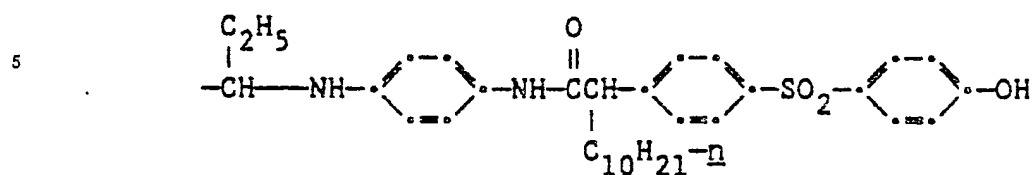


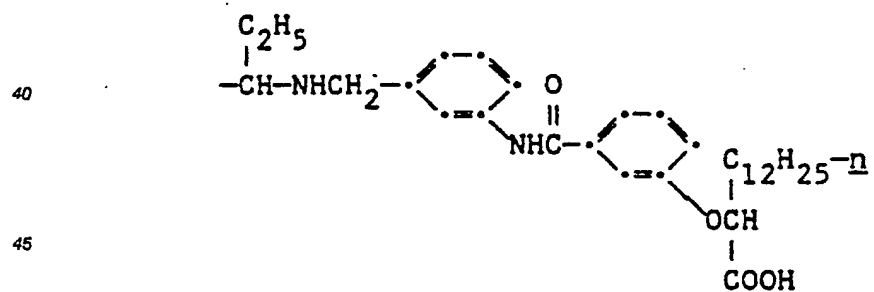
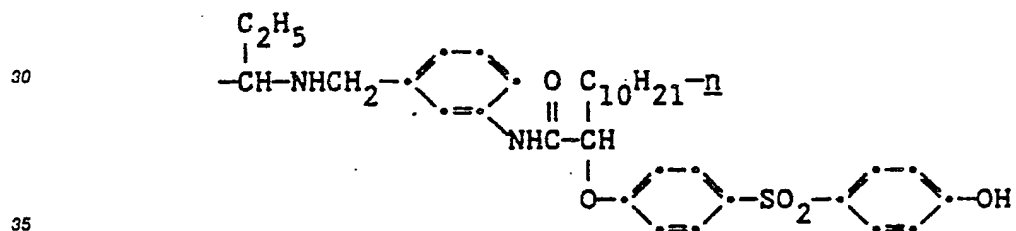
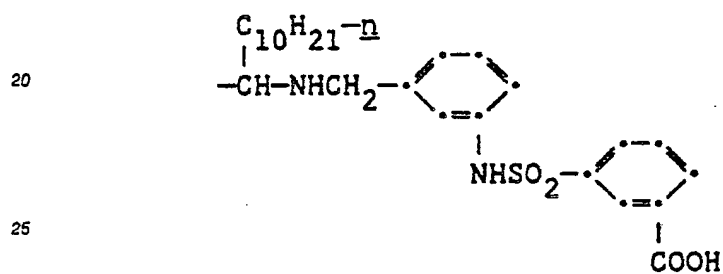
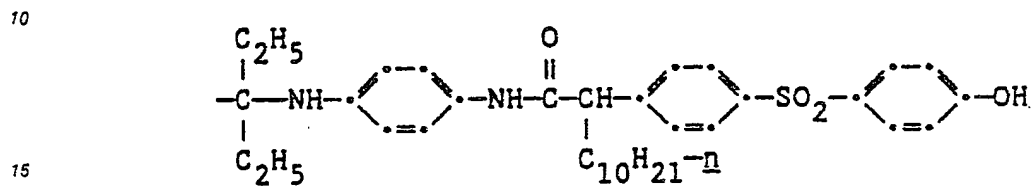
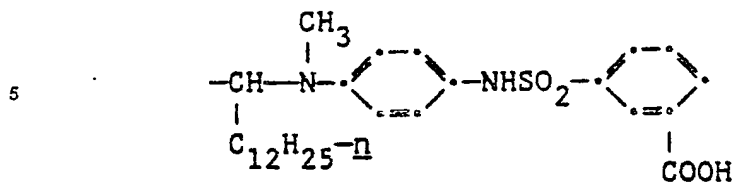
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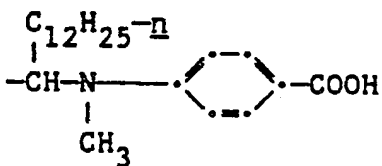
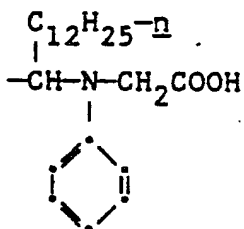
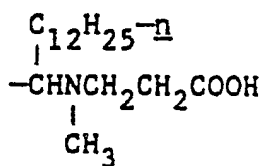
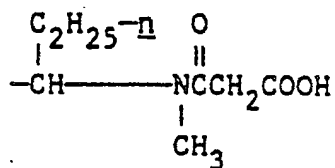
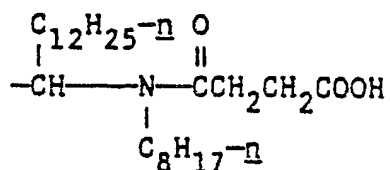
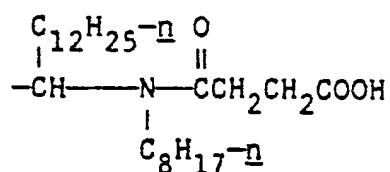
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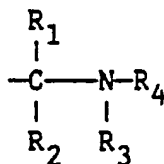


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Particularly useful pyrazoloazole couplers are 1H-pyrazolo[3,2-c]-s-triazole couplers, according to the invention, and contain the group



as described in the 3-position. Such couplers preferably contain in the 6-position an aryl group, particularly a phenyl or naphthyl group, which can be unsubstituted or optionally substituted, or an alkyl unsubstituted or substituted group, such as alkyl containing 1 to 40 carbon atoms.

A pyrazoloazole coupler, particularly a pyrazolotriazole coupler, typically contains a group which can aid solubility or diffusion resistance and produces a dye hue of desired hue upon reaction of the coupler with an oxidized color developing agent. For example, 1H-pyrazolo[3,2-c]-s-triazole couplers, according to the invention typically contain such a group in the 6-position.

5 A pyrazolotriazole coupler, for example, typically contains on the pyrazolotriazole nucleus, for instance in the 6-position of a 1H-pyrazolo[3,2-c]-s-triazole, a substituent that does not adversely affect the coupler, such as one of the following groups: alkyl, such as alkyl containing 1 to 30 carbon atoms, for example, methyl, ethyl, propyl, n-butyl, t-butyl, octyl and eicosyl; aryl, such as aryl containing 6 to 30 carbon atoms, for example, phenyl, naphthyl and mesityl. Such groups also include amino; acylamino, such as acylamino  
10 containing 2 to 30 carbon atoms, for example, acetamido, benzamido and stearamido; ureido; carboxy; alkanesulfonyl, such as ethanesulfonyl and butanesulfonyl; cyano (-CN); carbamyl, such as methyl carbamyl and hexyl carbamyl; sulfamyl, such as dioctyl sulfamyl and methyloctadecyl sulfamyl; sulfonamido; carboxamido; cycloalkyl, such as cyclohexyl and cyclopentyl; alkoxy, such as alkoxy containing 1 to 30 carbon atoms, for example, methoxy, i-butoxy and dodecyloxy; alkoxycarbonyl, such as ethoxycarbonyl and  
15 dodecyloxycarbonyl; aryloxycarbonyl, such as phenoxycarbonyl; alkylthio, such as alkylthio containing 1 to 30 carbon atoms, for example, methylthio and i-butylthio; aryloxy, such as aryloxy containing 6 to 30 carbon atoms, for example, phenoxy and naphthoxy; arylthio, such as arylthio containing 6 to 30 carbon atoms, for example, phenylthio; and heterocyclic groups, such as comprised of atoms selected from the group consisting of carbon, oxygen, nitrogen and sulfur atoms necessary to complete a 5-or 6-member  
20 heterocyclic ring, such as pyridyl, benzoxazolyl, furyl and thienyl. These groups on the pyrazolotriazole coupler are unsubstituted or optionally substituted with groups that do not adversely affect the desired properties of the pyrazolotriazole coupler. Examples of useful substituents can include ballast groups and coupler moieties known to be useful in the photographic art, or alkyl, such as alkyl containing 1 to 4 carbon atoms, for example, methyl, ethyl and t-butyl.

25 An example of such a group that is useful in the 6-position of a 1H-pyrazolo[3,2-c]-s-triazole coupler is a tertiary carbon group:



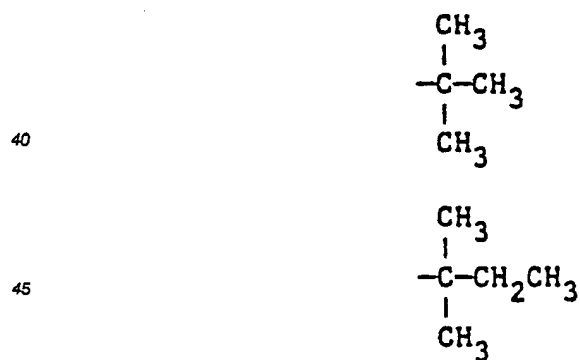
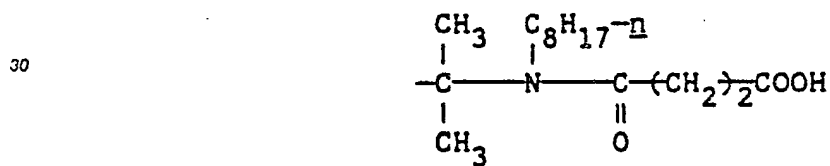
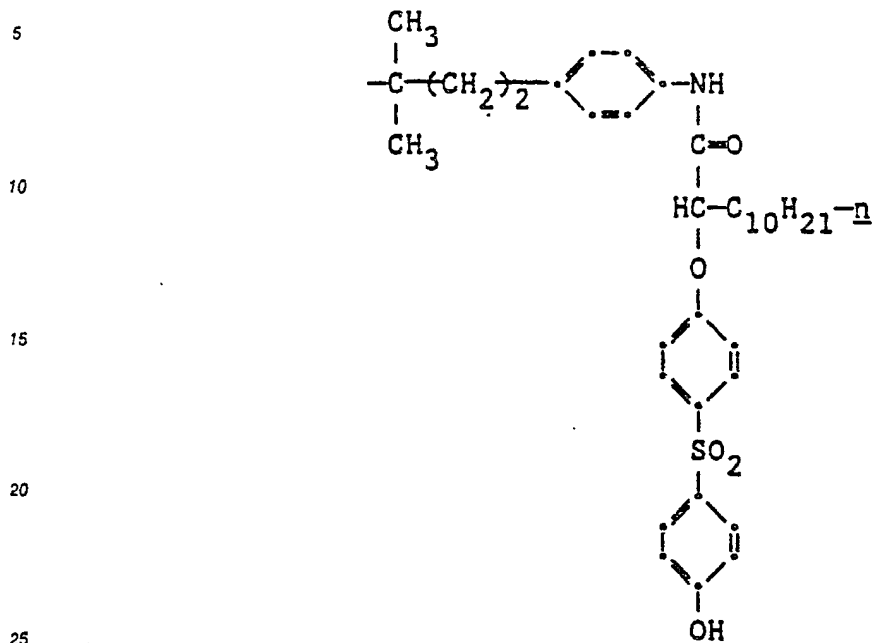
35 wherein:

$R_{18}$ ,  $R_{19}$ , and  $R_{20}$  are individually substituents that do not adversely affect the coupler, such as halogen, for example, chlorine bromine and fluorine; alkyl, such as alkyl containing 1 to 30 carbon atoms, for example, methyl, ethyl, propyl, butyl, pentyl, ethylhexyl and eicosyl; cycloalkyl, such as cyclohexyl and cyclopentyl; alkoxy, such as alkoxy containing 1 to 30 carbon atoms, for example, methoxy, ethoxy, butoxy  
40 and dodecyloxy; carboxy; amino, such as dioctylamino, dimethylamino and dodecylamino; carbonamido, such as acylamino containing 1 to 30 carbon atoms, for example, acetamido, stearamido, ureido and benzamido; alkylthio, such as alkylthio containing 1 to 30 carbon atoms, for example, methylthio, ethylthio, propylthio and dodecylthio; aryl, such as aryl containing 6 to 30 carbon atoms, for example, phenyl, naphthyl and mesityl; aryloxy, such as aryloxy containing 6 to 30 carbon atoms, for example, phenoxy and  
45 naphthoxy; cyano; nitro; sulfonamido; sulfamyl; carboxamido; alkylthio, such as ethylthio and butylthio; arylthio, such as arylthio containing 6 to 30 carbon atoms, for example, phenylthio and naphthylthio; or a heterocyclic group, such as a heterocyclic group comprised of atoms selected from carbon, oxygen, nitrogen and sulfur atoms necessary to complete a 5-or 6-member ring, for example, pyrrolyl, oxazolyl and pyridyl. Optionally, in such a tertiary group,  $R_{18}$  can form with one of  $R_{19}$  and  $R_{20}$  a heterocyclic ring, such  
50 as a heterocyclic ring comprised of atoms selected from carbon, nitrogen, oxygen and sulfur atoms necessary to complete a 5-or 6-member heterocyclic ring, for example, oxazole, pyridine, pyrrole and thiophene; or,  $R_{18}$  can form with one of  $R_{19}$  and  $R_{20}$  a carbocyclic ring, such as cyclohexyl or norbornyl; or  $R_{18}$ ,  $R_{19}$  and  $R_{20}$  can comprise the carbon and hydrogen atoms necessary to complete a ring, such as an adamantyl ring.

55 The groups  $R_{18}$ ,  $R_{19}$  and  $R_{20}$  are unsubstituted or optionally substituted with groups that do not adversely affect the desired properties of the pyrazolotriazole coupler. The groups can be optionally substituted with, for example, alkyl, such as alkyl containing 1 to 20 carbon atoms, for example, methyl, ethyl, propyl and butyl; aryl, such as phenyl and naphthyl; or phenolic, carboxylic acid and heterocyclic

substituent groups. Substituents can include ballast groups and coupler moieties which are known to be useful in the photographic art.

Examples of useful tertiary carbon groups are



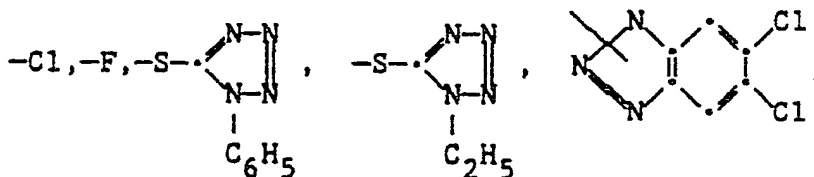
The pyrazoloazole couplers contain in the coupling position, that is the 7-position in a 1H-pyrazolo[3,2-

c)-s-triazole coupler, hydrogen or a coupling-off group, also known as a leaving group.

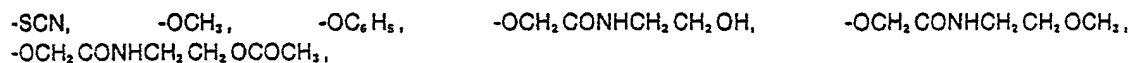
Coupling-off group, defined by Q herein, are well known to those skilled in the art. Such groups can determine the equivalency of the coupler, can modify the reactivity of the coupler, or can advantageously affect the layer in which the coupler is coated or other layers in the element by performing, after release  
 5 from the coupler, such functions as development inhibition, development accelerations, bleach inhibition, bleach acceleration, color correction and the like. Representative classes of coupling-off groups include halogen, particularly chlorine, bromine and fluorine, alkoxy, carbonamido, imido, aryloxy, particularly substituted phenoxy, heterocycloxy, sulfonyloxy, acyloxy, heterocyclyl, thiocyno, alkylthio, arylthio, particularly substituted phenylthio, heterocyclylthio, sulfonamido, phosphonyloxy and arylazo. They are described  
 10 in, for example, U.S. Patents 2,355,169; 3,227,551; 3,432,521; 3,476,563; 3,617,291; 3,880,661; 4,052,212 and 4,134,766; and in U.K. patents and published application numbers 1,466,728; 1,531,927; 1,533,039; 2,006,755A and 2,017,704A.

Examples of specific coupling-off groups are

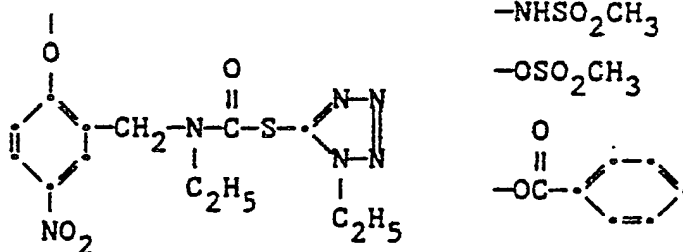
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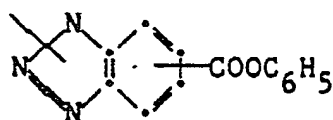
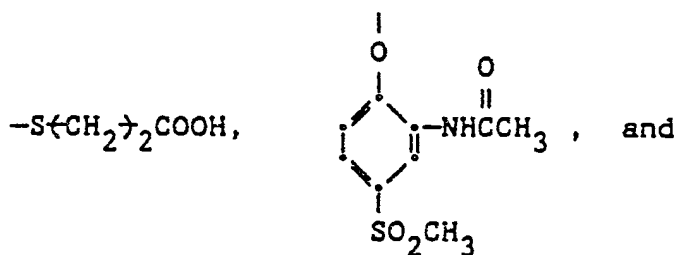
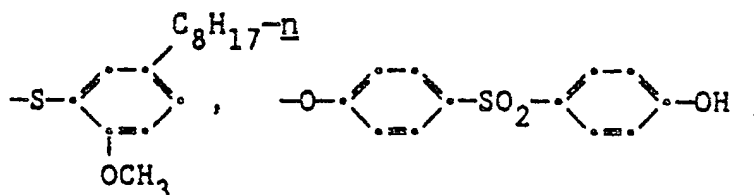
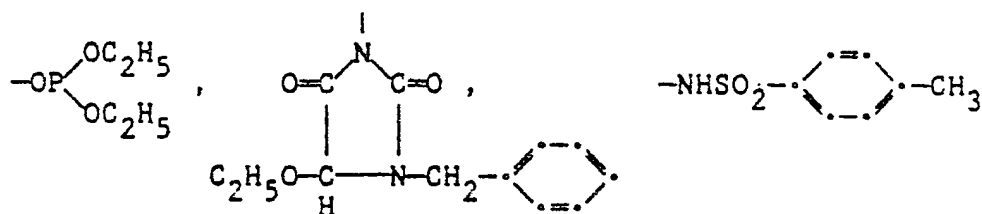
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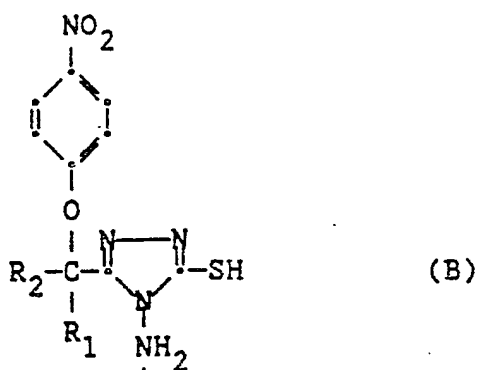
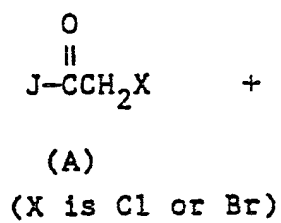
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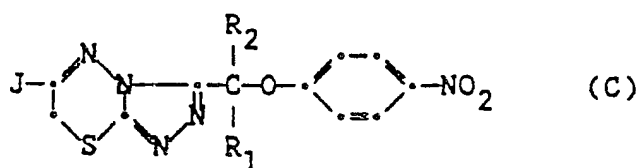




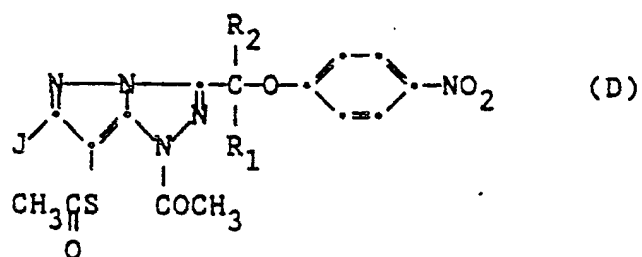
A ballast group as described is an organic radical of such size and configuration as to confer on the coupler molecule sufficient bulk to render the coupler substantially non-diffusible from the layer in which it is coated in a photographic element. Couplers of the invention can contain ballast groups, or be bonded to polymeric chains through one or more of the groups described herein. For example, one or more coupler moieties can be attached to the same ballast group. Representative ballast groups include substituted or unsubstituted alkyl or aryl groups containing 8 to 32 carbon atoms. Representative substituents include alkyl, aryl, alkoxy, aryloxy, alkylthio, arylthio, hydroxy, halogen, alkoxycarbonyl, aryloxycarbonyl, carboxy, acyl, acyloxy, carbonamido, carbamoyl, alkylsulfoxide, arylsulfoxide, alkanesulfonyl, arenesulfonyl, amino, anilino, sulfonamido and sulfamoyl groups wherein the alkyl and aryl substituents and the alkyl and aryl portions of the alkoxy, aryloxy, alkylthio, arylthio, alkoxycarbonyl, aryloxycarbonyl, acyl, acyloxy, carbonamido, carbamoyl, alkanesulfonyl, arenesulfonyl, sulfonamido and sulfamoyl substituents containing 1 to 30 carbon atoms and 6 to 30 carbon atoms, respectively, and can be further substituted with such substituents.

Pyrazoloazole couplers according to the invention are prepared by the general method of synthesis described in Research Disclosure, August 1974, Item No. 12443 published by Kenneth Mason Publications, Ltd., The Old Harbourmaster's, 8 North Street, Emsworth, Hampshire PO10 7DD, England and U.S. Patent 4,540,654. An illustrative synthesis scheme I is as follows:

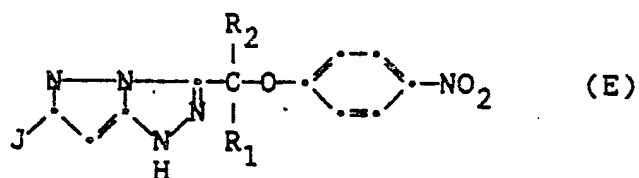




10  
reflux with  
acetic anhydride



25  
reflux with HCl  
and acetic acid



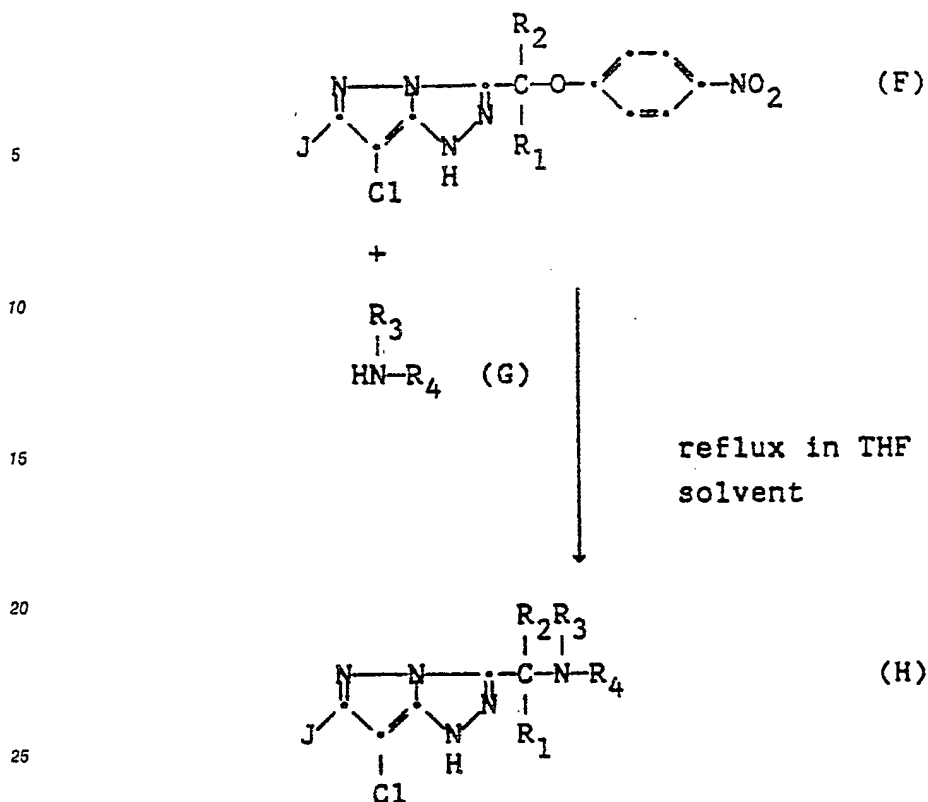
35  
(remove elemental  
sulfur formed)

40  
(add optional  
coupling-off group,  
such as chlorine by  
reaction with  
N-chlorosuccinimide)

45

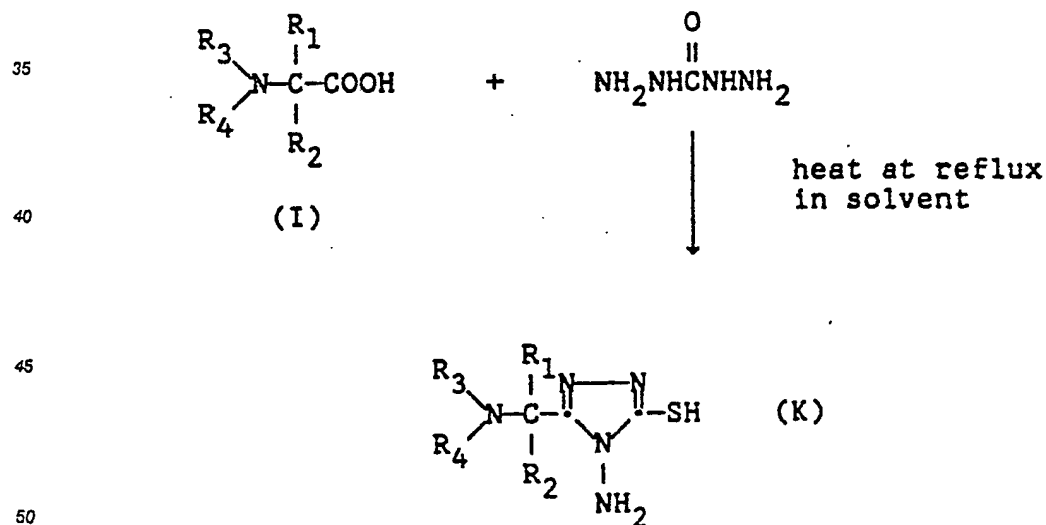
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55



wherein R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub>, R<sub>4</sub> and J are as defined.

30 An optional method for preparing an intermediate (K) for preparing 1H-pyrazolo[3,2-c]-s-triazole couplers according to the invention is illustrated in the following method:

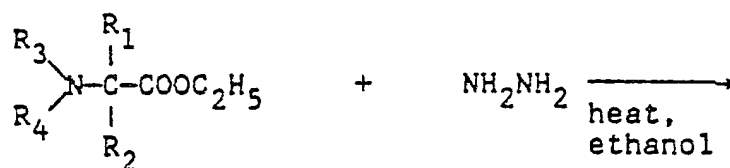


wherein R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub> and R<sub>4</sub> are as described.

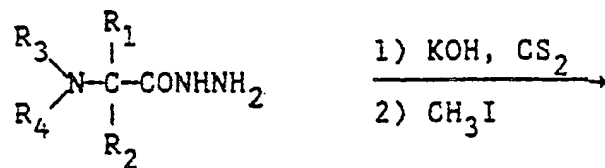
55 This intermediate (K) can be useful to form pyrazolotriazole (H) without the need for formation of a p-nitrophenoxy group as in intermediate (B).

Another optional method for preparing an intermediate (K) for preparing pyrazolotriazoles according to the invention is illustrated in the following sequence:

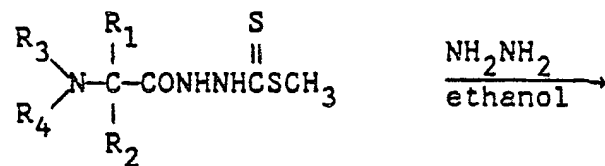
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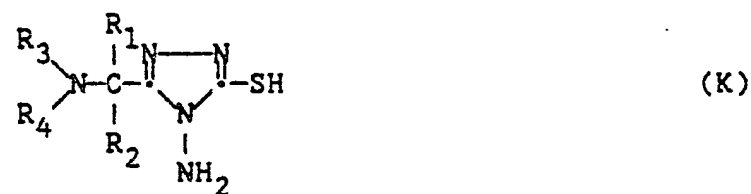
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25

wherein  $R_1$ ,  $R_2$ ,  $R_3$  and  $R_4$  are as described.

30 Illustrative examples of the preparation of pyrazoloazole couplers according to the invention are as follows:

(Et herein means  $-C_2H_5$ .)

( $Ac_2O$  herein means acetic anhydride.)

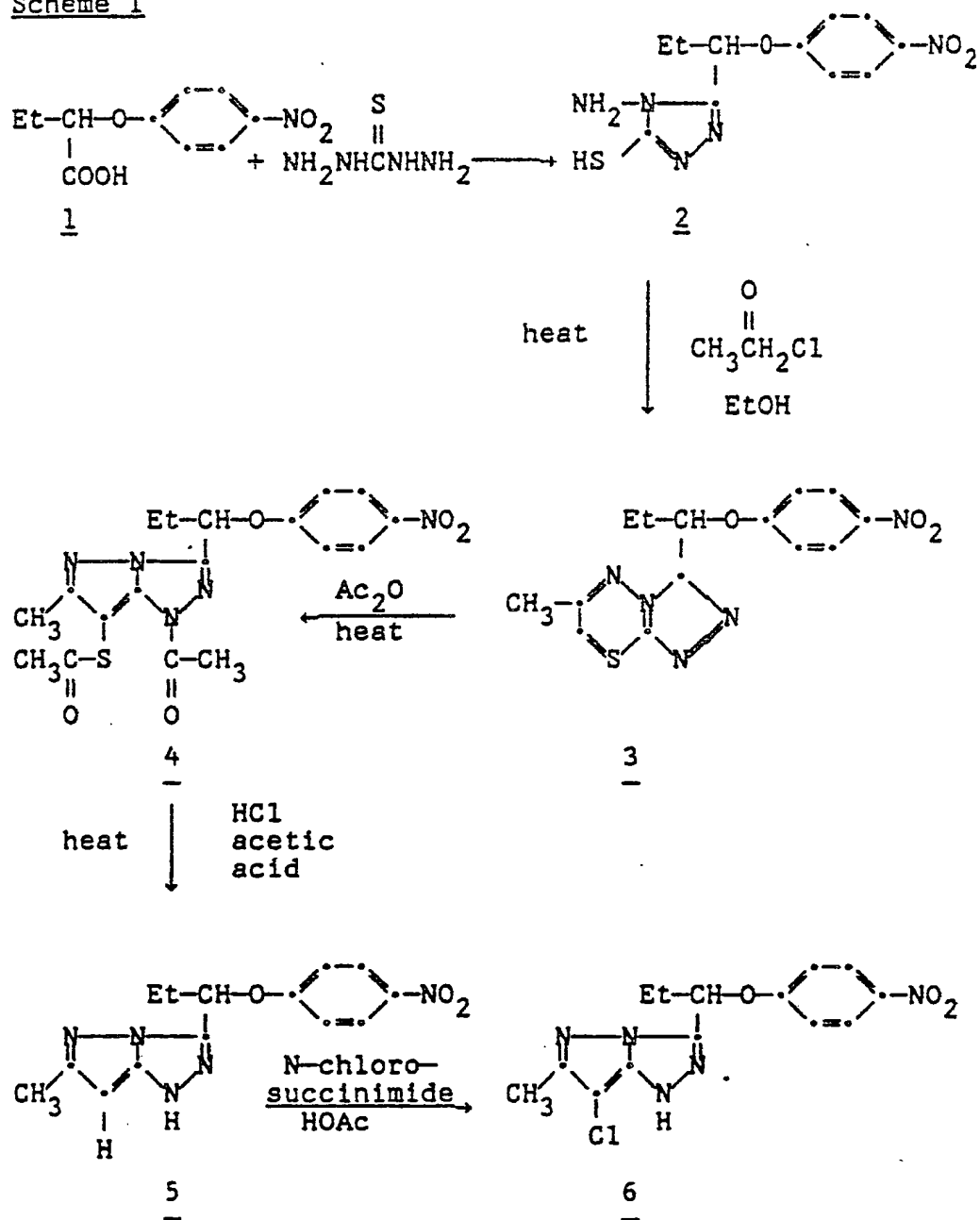
35

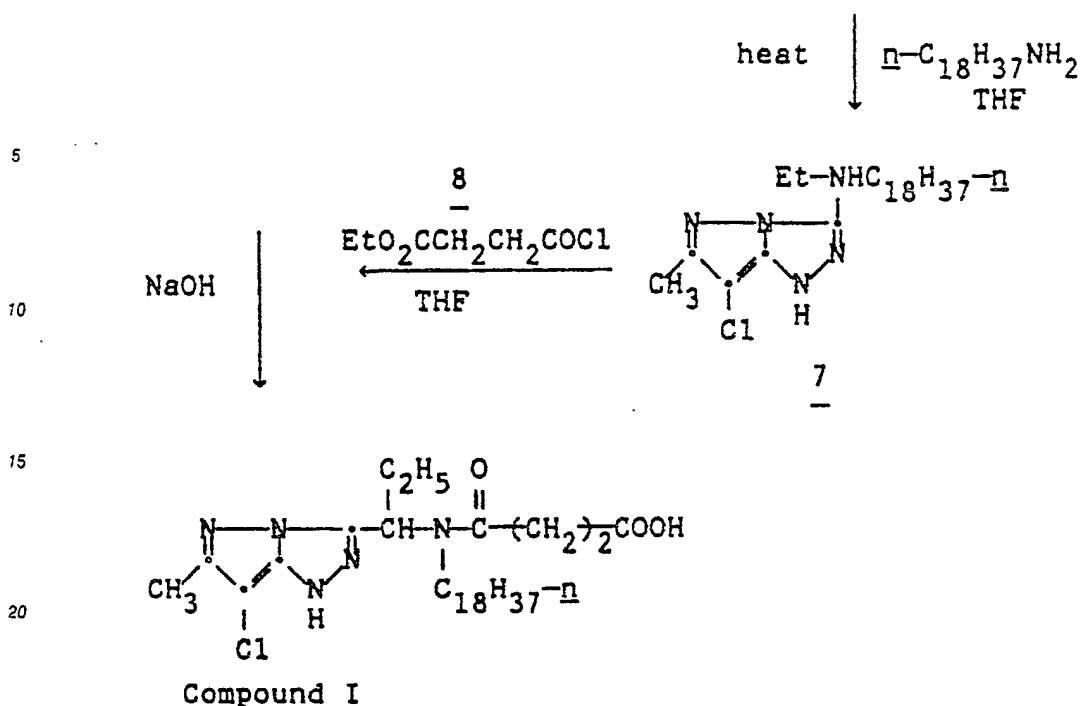
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45

50

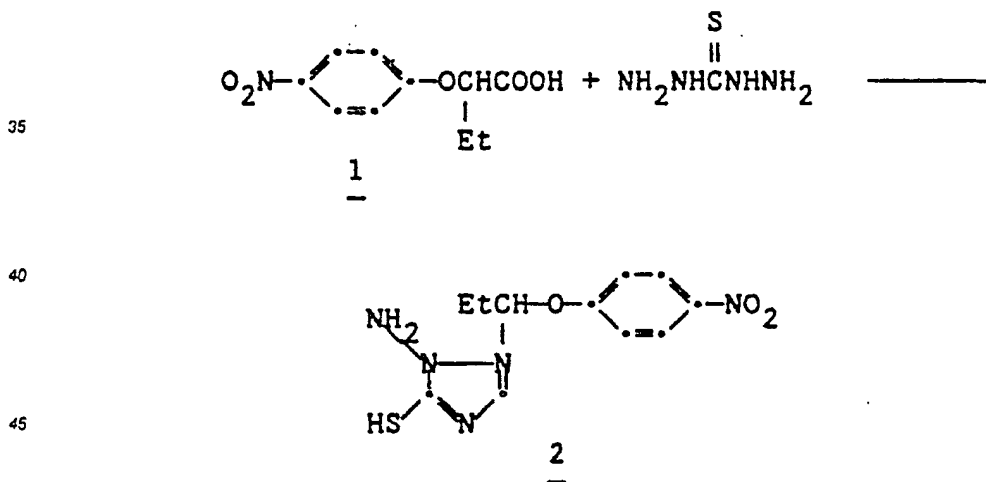
55

Synthesis Example AScheme I



# Synthesis Example A

30 (Preparation of Intermediate 2)



50 10.0 gm (.044 mol) acid 1 and 5.2 gm (.049 mol) thiocarbonylhydrazide were mixed intimately and heated to 130°C for three hours. EtOH was added to the hot mixture and the mixture allowed to cool with stirring overnight. The desired intermediate was a white solid (8.1 gm) (62%). The intermediate was filtered and air dried. (NMR and nmr herein mean nuclear magnetic resonance.)

NMR analysis:

55 H nmr:  $\delta$  1.0 (t, 3H); 2.22 (q, 2H); 5.6-5.9 (s, m, 3H); 7.25 (d, 2H); 8.25 (d, 2H).

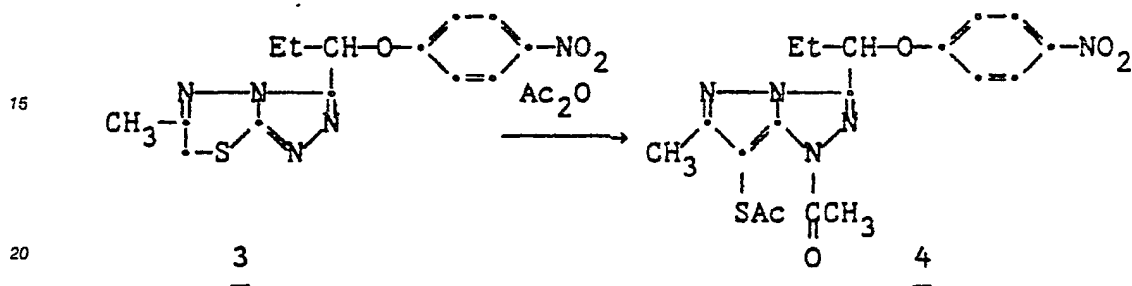
## (Preparation of Intermediate 3)

10 grams (.034 mol) of 2 were refluxed 3 hours with 3.6 ml (.045 mol) chloroacetone, cooled, and impregnated on silica gel. The product was chromatographed to obtain 9.2 gm of yellow glass. This was  
 5 triturated overnight with ether to give 8.2 gm (73%) light tan solid 3.

NMR Analysis:

<sup>1</sup>H NMR:  $\delta$  1.1 (t,3H); 2.0-22.5 (s,q, 5H); 3.65 (s,2H), 7.68 (t,1H); 7.15 (d,2H) 8.15 (d,2H)

## 10 (Preparation of Intermediate 4)



25 8 gm (.024 mol) of 3 were refluxed overnight in acetic anhydride. The product was cooled and added to a mixture of water and ether and then stirred 4 hours. The organic layer washed with water, dried over magnesium sulfate, and evaporated to a brown viscous oil, 8.3 gm.

NMR Analysis:

<sup>1</sup>H NMR:  $\delta$  1.2 (t,3H); 2.1-2.7 (s,s,s,m,1H); 5.5(t,1H); 7.05 (d,2H) 8.05 (d,2H).

30

## (Preparation of Intermediate 5)

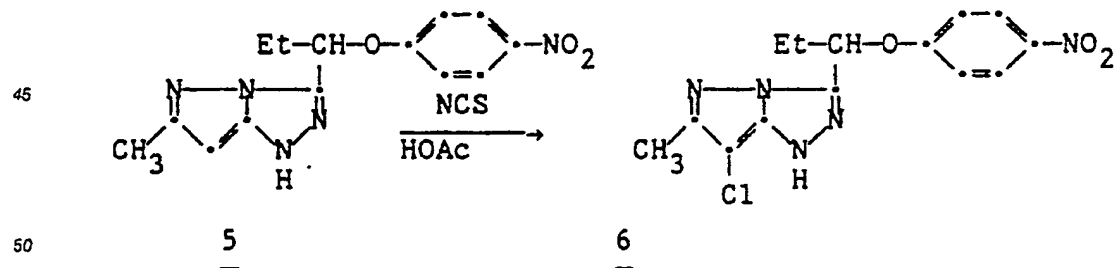
35 5.0 gm of 4 was refluxed one hour with 50 ml CONC HCl and 50 ml glacial acetic acid. The product was cooled and added to water. Ethyl acetate was added and the organic layer washed, dried ( $MgSO_4$ ), and evaporated to 3.4 gm oil. The product was triturated with  $CH_3CN$  to yield 1.7 gm (47%) white solid 5.

NMR Analysis:

<sup>1</sup>H NMR:  $\delta$  0.9 (t,3H); 1.0-1.8 (m,2H); 5.58 (5,1H); 5.62 (6,1H); 7.13 (d,2H); 8.16 (d,2H).

40

## (Preparation of Intermediate 6)



55 .86 gm (.0064 mol) N-chlorosuccinimide added to 2.0 gm (.0066 mol) 5 dissolved in 10 ml acetic acid and stirred one hour at room temperature. The mixture added to water, ethyl acetate added and the organic layer washed with water, dried ( $MgSO_4$ ), and evaporated to light yellow solid, 2.1 gm (98%).

NMR Analysis:

<sup>1</sup>H NMR:  $\delta$  1.05 (t,3H); 2.15-2.65 (s,m,5H); 5.55 (t,1H); 7.1 (d,2H); 8.15 (d,2H); 9.55 (s,1H).



(Preparation of Intermediate 7)

A mixture of 5.0 gm (.015 mol) of 6 and 4.0 gm (.015 mol, octadecylamine were refluxed one hour in 200 ml THF. The product was cooled and added to water. Ether was added and the organic layer washed with water, twice with saturated  $\text{Na}_2\text{CO}_3$  and evaporated to 7.5 gm brown oil. The brown oil was chromatographed on silica gel to yield 4.3 gm (62%) light brown oil 7.

Analysis:

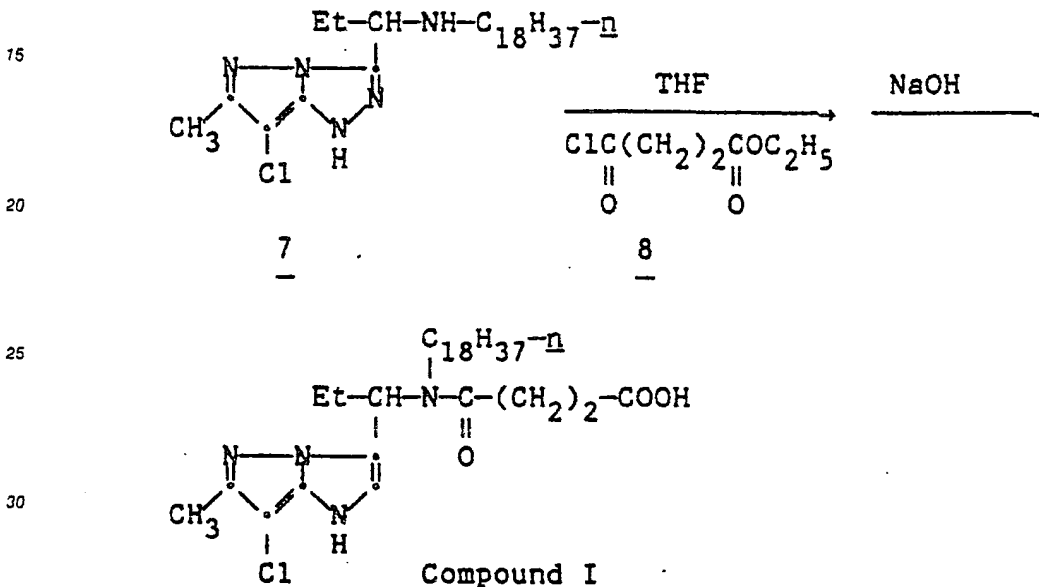
Found

C: 67.2; H:10.1; N: 15.1.

10 Theo

C: 67.0; H: 10.4; N: 15.0.

NMR Analysis: ( $\text{CDCl}_3$ ):  $\delta$  .90 (m,6H), 1.0-1.6 (m,32H), 1.98-2.7 (m,7H), 4.15 (t,1H).



35 2.1 gm (.0045 mol) 7 was refluxed with 1.68 gm (.0102 mol) 8 in 30 ml THF for 10 minutes. The product was added to ether and water and the organic layer washed, dried and evaporated. The residue dissolved in 20 ml THF and 20 ml methanol and 1 ml of a 50% solution of NaOH in water added. The product was stirred at ambient temperature for one half hour and then added to dilute HCl. Ether was added and the organic layer washed, dried and evaporated to dark oil. The product was chromatographed on silica gel to obtain 1.8 gm oil.

Analysis:

Found:

N: 11.8; C: 63.1; H: 9.6.

Theo:

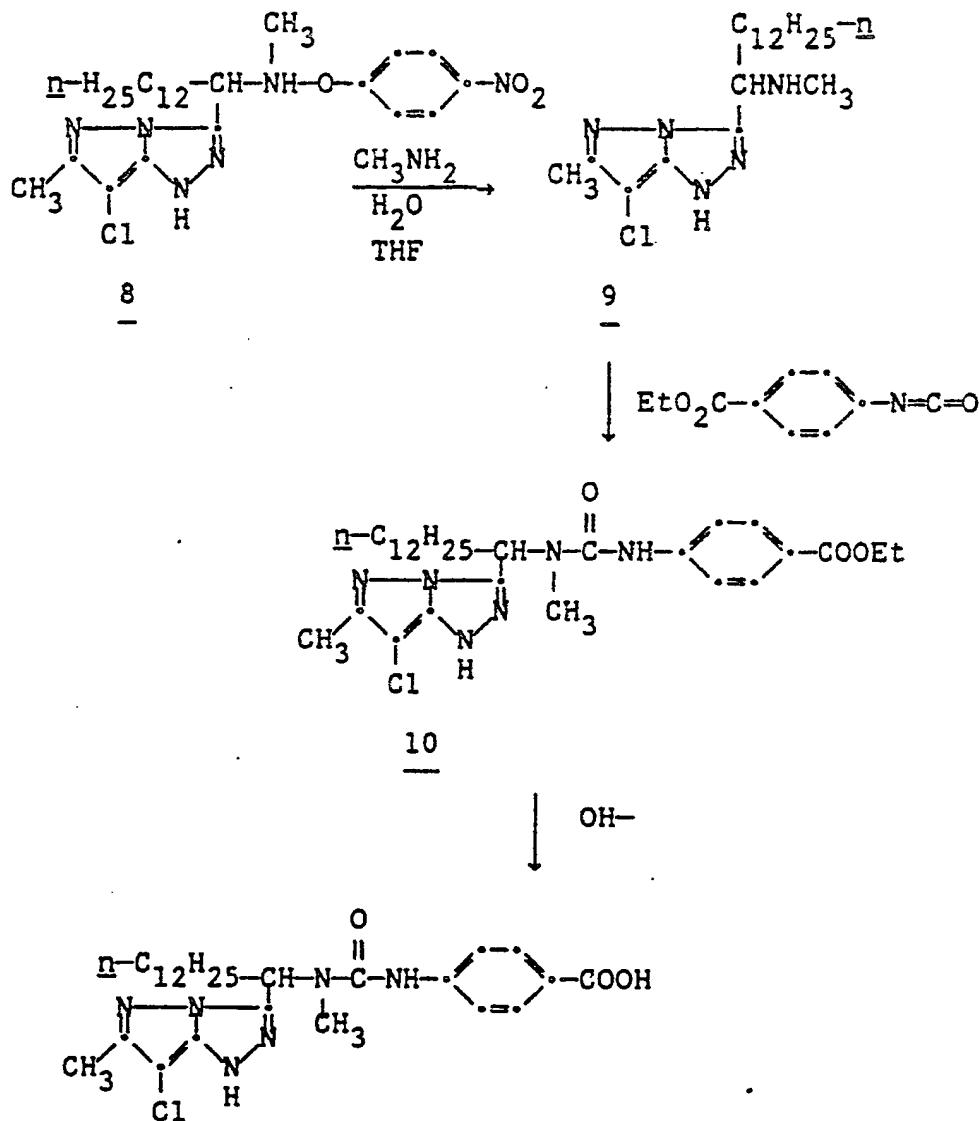
45 N:12.4; C: 63.6; H: 9.3.

50

55

## Synthesis Example B

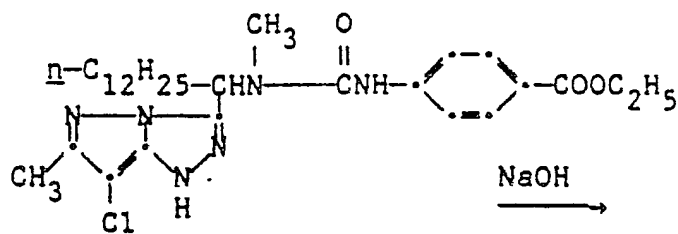
## Scheme II



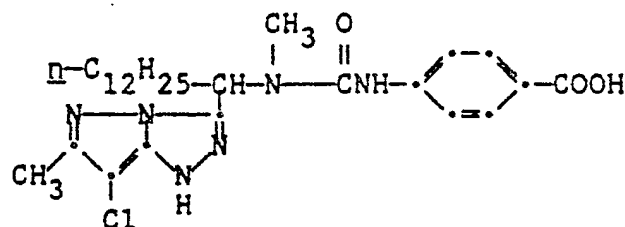
## Compound II

16 ml of a solution of methylamine in water, (40%), was added with stirring to 20 gm of compound 8 - (obtained in the same manner as 6) and stirred at room temperature for one half hour. Water and ether were added and the organic layer washed with water, then  $\text{Na}_2\text{CO}_3$  solution. Water and dried ( $\text{MgSO}_4$ ) and evaporated to solid. The product was triturated with ether to obtain 6.6 gm white solid 9.

5.4 gm (.015 mol) 9 and 2.64 gm (.015 mol) of isocyanate were refluxed in 20 ml THF for 5 minutes. After cooling, water and ethyl acetate were added. The organic layer was washed, dried, and evaporated to a tan solid. The product was triturated with acetonitrile to give 7.6 gm off-white solid 10.



NaOH



Compound II

7.5 gm 10 was dissolved in 100 ml THF and 100 ml MeOH and 30 ml of a 50% solution of sodium hydroxide were added. After stirring one hour the reaction mixture was added to dilute HCl. Ethyl acetate was added and the organic layer washed, dried, and evaporated. The product was crystallized from acetonitrile to give 6.0 gm white solid.

MP 178.5-180°C

Analysis

Found:

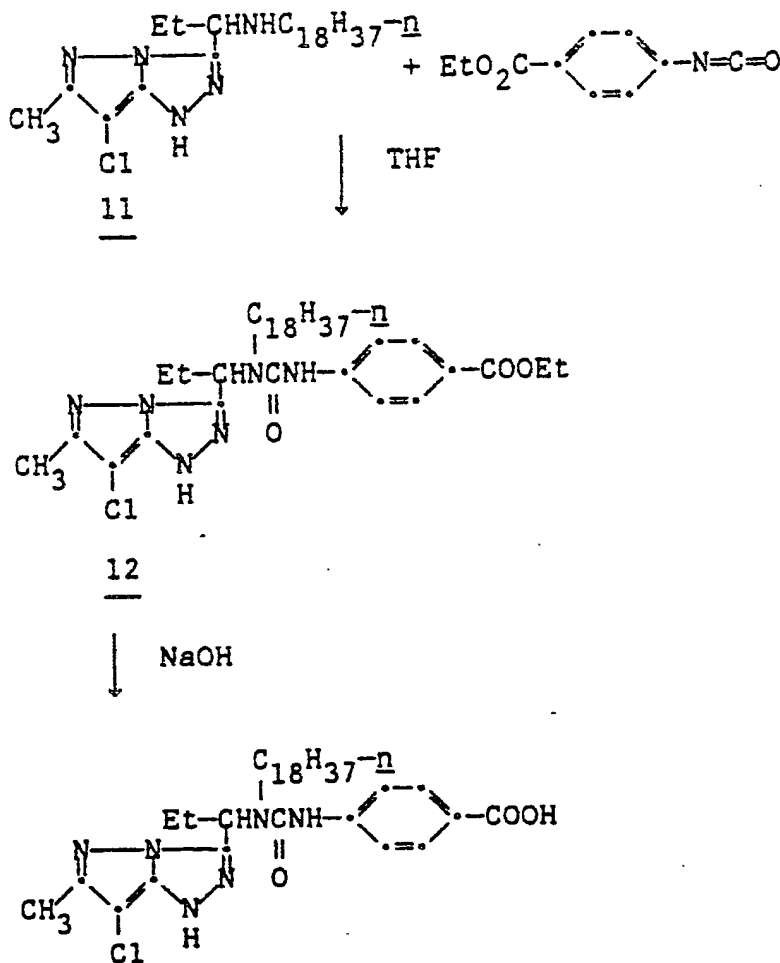
N: 15.9; C: 61.0; H: 7.5

Theoretical :

N: 16.0; C: 60.2; H: 7.6

## Synthesis Example C

Scheme III



40 5.0 gm (.011 mol) of 11 and 4.0 gm (.022 mol) of the isocyanate were refluxed one half hour in 20 ml THF. The product was cooled and added to water and ethyl acetate. The organic layer was washed, dried, and evaporated and chromatographed on silica gel. The product was triturated with  $\text{CH}_2\text{CN}$  to obtain 5.2 gm white solid 12.

45 50 grams of 12 were dissolved in a mixture of 100 ml THF and 100 ml MeOH and 20 ml of a 50% solution of NaOH in water added. After stirring at room temperature one hour the product was added to dilute HCl. Ethyl acetate was added and the organic layer washed, dried and evaporated. The solid was crystallized from acetonitrile to give 4.5 grams white solid Compound III.  
MP 178.5-180.0°C.

50 Analysis:

Found

N: 13.3; C: 65.2; H: 8.7;

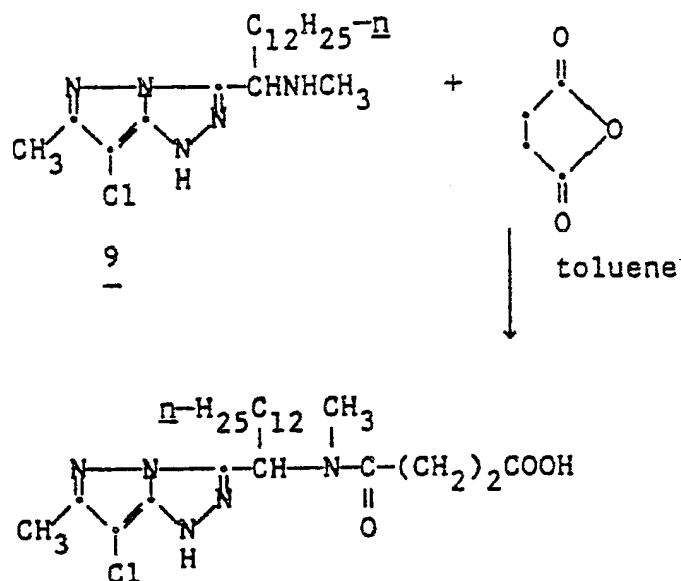
Theoretical

N: 13.4; C: 64.9; H 8.5

55

## Synthesis Example D

## Scheme IV



5.8 gm (.016 mol) of 9, preparation of which is described in Scheme II was refluxed 10 minutes with 1.7 gm (.017 mol) of succinic anhydride in 25 ml toluene and added to water. The organic layer was washed, dried and evaporated and chromatographed on silica gel. The product was triturated with acetonitrile to yield a white solid, 3.0 gm. mp 112-113°C.

Analysis:

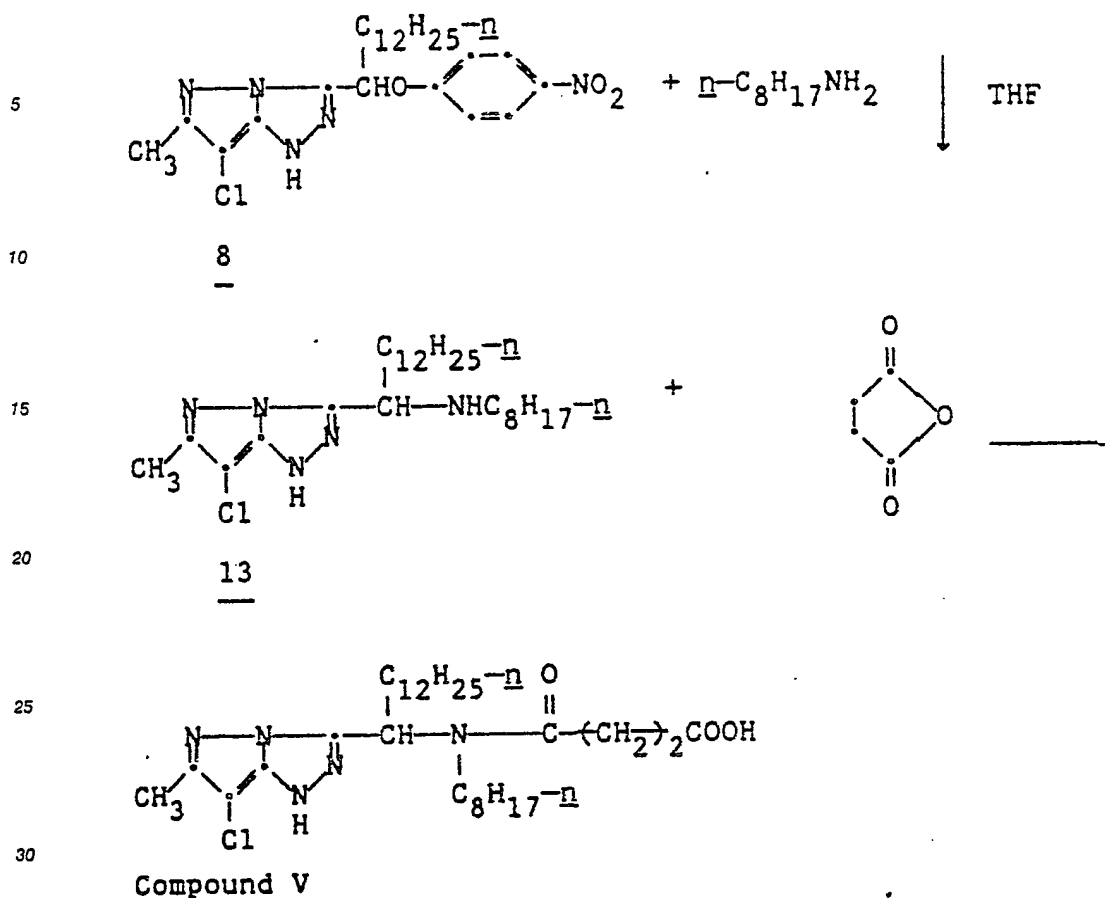
Found:

N: 14.9; C: 59.1; H: 7.9;

Theoretical:

N: 15.0; C: 59.0; H 8.2.

## Synthetic Example E



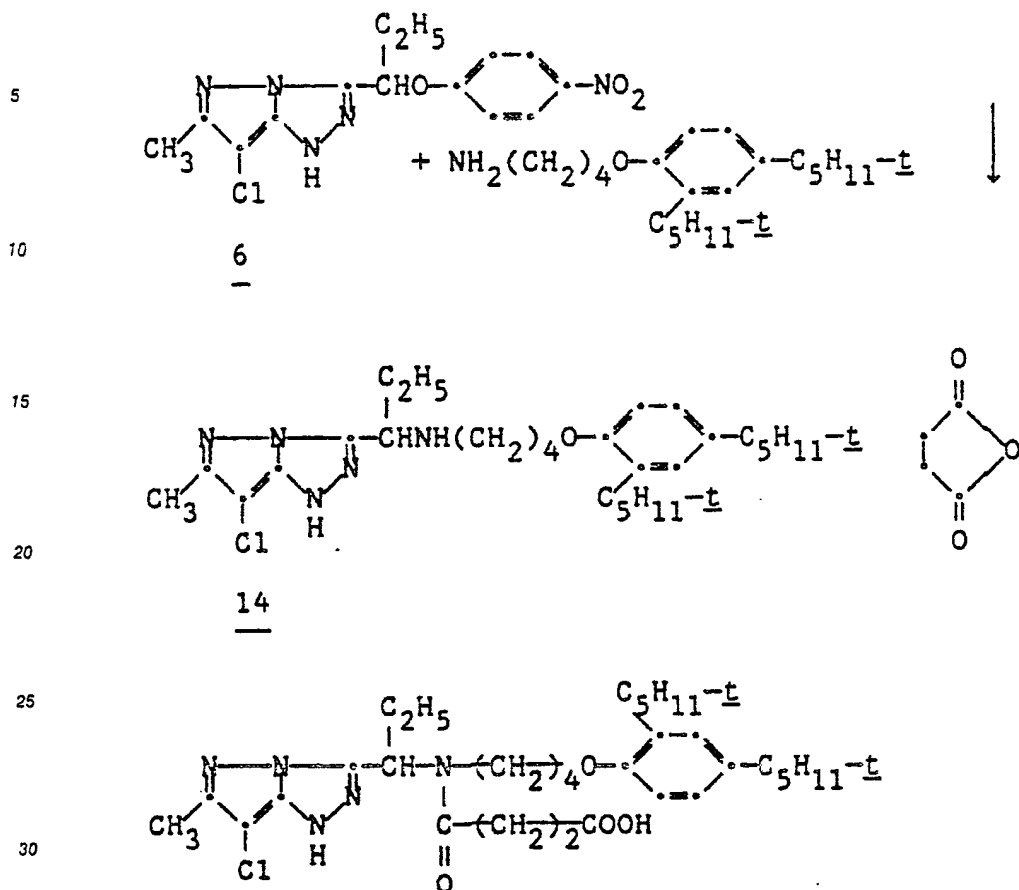
Compound V was made from 13 in the same way as Compound IV was made from 9 (see Scheme IV).

Compound 13 was made from 8 in the same way as 9 was made from 8.

NMR Analysis:

<sup>1</sup>H NMR:  $\delta$  8.9 (t, 3H); .9-1.6 (m, 40H); 1.9-2.5 (m, s, 5H); 2.6-2.9 (m, 4H); 3.15-3.50 (m, 2H); 5.3 and 6.0 (t, 1H).

## Synthetic Example F



35

Compound 14 was prepared from 6 in the same manner as 7 was prepared from 6 and subsequently converted into Compound VI in the same fashion as 13 was converted to Compound V.

NMR Analysis:

40 <sup>1</sup>H NMR: δ .65 (t,3H); .75 (t,3H); 1.3-1.5 (m,12H); 1.6-2.0 (m,8H); 2.37 (s,3H); 2.39-2.55 (m,2H); 2.7-3.0 (m,4); 3.3-3.6 (m,2H); 3.85-4.0 (m,2H); 5.3 + 6.0 (t,t,1H); 6.83 (m,1H); 7.18 (m,1H); 7.25 (s,1H).

Analysis:

Found:

C: 63.9; H: 8.1; N: 11.6

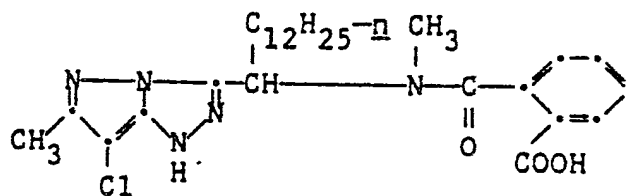
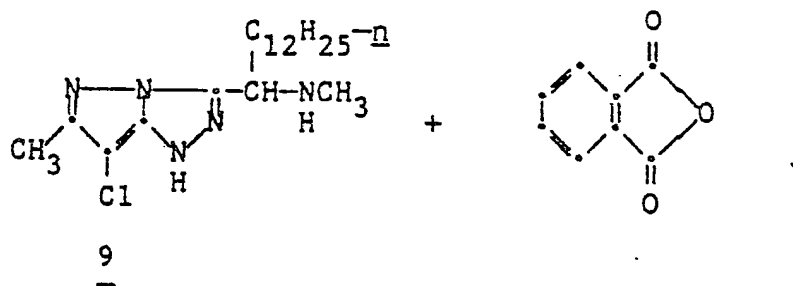
Theoretical:

45 C: 63.8; H: 8.0; N: 11.6

50

55

## Synthetic Example G



Compound VII

25 Compound VII was prepared from 9 in the same way as IV was prepared from 9.

Analysis:

Found:

C: 63.2; H: 7.5; N: 13.8

Theoretical:

C: 62.8; H: 7.4; N: 13.6

30 <sup>1</sup>H NMR: .9 (t,3H); 1.0-1.6 (m,23H); 2.0-2.4 (m,s,sh); 2.8 (s,3H); 5.0 + 2.6.3 (m;m,1H); 7.35-7.8 (m,3H); 8.19 (t,1H).

The following compounds can be prepared in a similar manner:

35

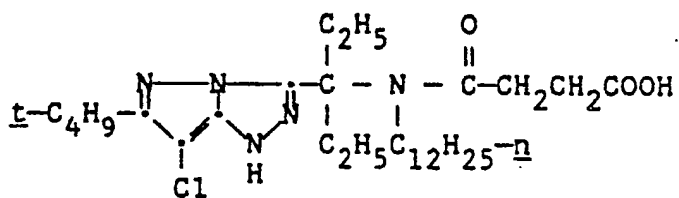
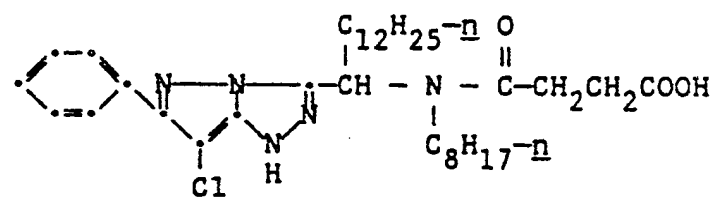
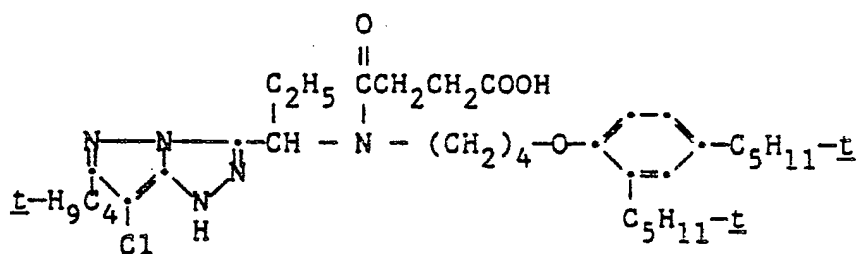
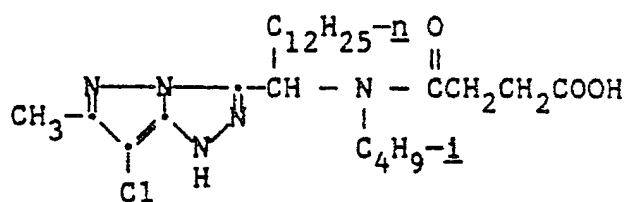
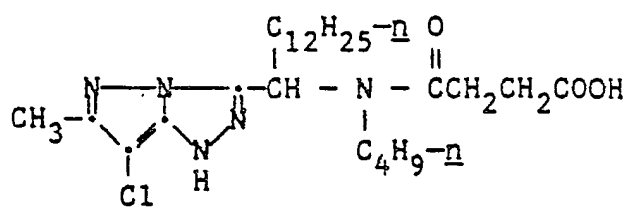
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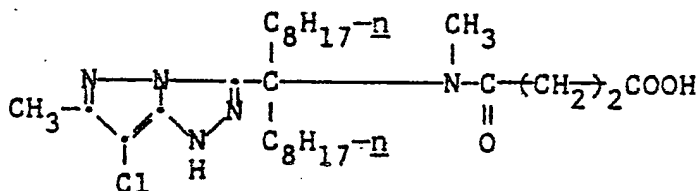
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The photographic elements of this invention or individual layers thereof, can contain, for example, brighteners (see Research Disclosure Section V), antifoggants and stabilizers (See Research Disclosure Section VI), antistain agents and image dye stabilizer (see Research Disclosure Section VII, paragraphs I and J), light absorbing and scattering materials) see Research Disclosure Section VIII), hardeners (see Research Disclosure Section XI), plasticizers and lubricants (see Research Disclosure Section XIII), matting agents (see Research Disclosure Section XVI) and development modifiers (see Research Disclosure Section XXI) colored masking couplers, bleach accelerators, inhibitors and competing couplers.

The photographic elements can be coated on a variety of supports as described in Research Disclosure Section XVII and the references described therein.

Photographic elements can be exposed to actinic radiation, typically in the visible region of the spectrum, to form a latent image as described in Research Disclosure Section XVIII and then processed to form a visible dye image as described in Research Disclosure Section XIX. Processing to form a visible dye image includes the step of contacting the element with a color developing agent to reduce developable silver halide and oxidize the color developing agent. Oxidized color developing agent in turn reacts with the coupler to yield a dye.

Preferred color developing agents are p-phenylene diamines. Especially preferred are

4-amino-3-methyl-N,N-diethylaniline hydrochloride,

4-amino-3-methyl-N-ethyl-N-β-(methanesulfonamido)ethylaniline sulfate hydrate,

4-amino-3-methyl-N-ethyl-N-β-hydroxyethylaniline sulfate,

4-amino-3-β-(methanesulfonamido)ethyl-N,N-diethylaniline hydrochloride and

4-amino-N-ethyl-N-(2-methoxyethyl)-m-toluidine-di-p-toluenesulfonic acid.

With negative working silver halide this processing step leads to a negative image. To obtain a positive (or reversal) image, this step can be preceded by development with a non-chromogenic developing agent to develop exposed silver halide, but not form dye, and then uniform fogging of the element to render unexposed silver halide developable. Alternatively, a direct positive emulsion can be employed to obtain a positive image.

Development is followed by the conventional steps of bleaching, fixing, or bleach-fixing, to remove silver and silver halide, washing, and drying.

The following examples further illustrate the invention.

#### Examples 1-37 -(Photographic Elements Comprising Pyrazoloazole Couplers of the Invention)

Photographic elements were prepared by coating a cellulose acetate-butyrate film support with a photosensitive layer containing a silver bromiodide emulsion at 0.91 gm Ag/m<sup>2</sup>, gelatin at 3.77 gm/m<sup>2</sup>, and one of the couplers designated in Table II dispersed in half its weight of tricresyl phosphate and coated at 1.62 mmol/m<sup>2</sup>. The photosensitive layer was overcoated with a layer containing gelatin at 1.08 gm/m<sup>2</sup> and bis-vinylsulfonylmethyl ether at 1.75 weight percent based on total gelatin.

Samples of each element were imagewise exposed through a graduated-density test object and processed at 40°C employing the following color developing solution, then stopped, bleached, fixed, washed, and dried to produce stepped magenta dye images.

K<sub>2</sub>SO<sub>3</sub> 2.0 gm

K<sub>2</sub>CO<sub>3</sub> 30.0 gm

KBr 1.25 gm

KI 0.6 gm

4-Amino-3-methyl-N-ethyl-N-β'-hydroxyethylaniline sulfate 3.55 gm

Water to 1.0 liter, pH 10.0

Densitometry of these images provided measures of maximum density (D<sub>max</sub>). Dye hues were obtained from spectrophotometric curves by measuring the maximum absorption peak (λ<sub>max</sub>) normalized to 1.0 density.

Described Compound No. III was used as the magenta dye-forming coupler of Example 1. Described Compound II was used as the magenta dye-forming coupler of Example 2. The results are given in the following Tables.

Table II

<u>Example No.</u>	<u>Compound No.</u>	<u><math>\lambda_{max}</math></u>	<u>D<sub>max</sub></u>	<u>D<sub>min</sub></u>
1	III	560	3.94	.21
2	II	567	3.98	.14
Comparison A	*C-8	555	3.03	.13

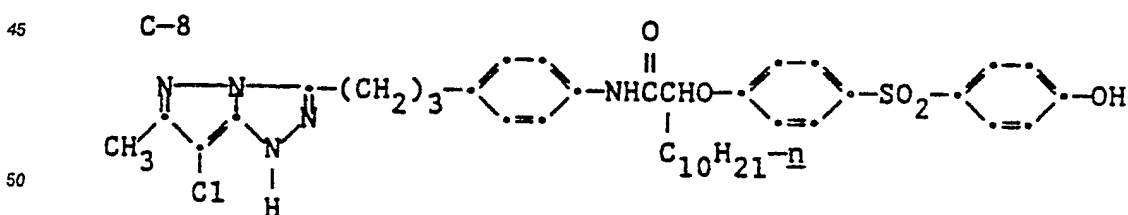
Table III

<u>Example No.</u>	<u>Compound No.</u>	<u><math>\lambda_{max}</math></u>	<u>D<sub>max</sub></u>	<u>D<sub>min</sub></u>
3	V	562	3.93	.15
Comparison A	*C-8	556	3.62	.14

Table IV

<u>Example No.</u>	<u>Compound No.</u>	<u><math>\lambda_{max}</math></u>	<u>D<sub>max</sub></u>	<u>D<sub>min</sub></u>
4	VI	561	4.05	.13
5	VII	563	3.87	.12
Comparison A	*C-8	556	3.32	.14

\*Structure of comparison coupler C-8:

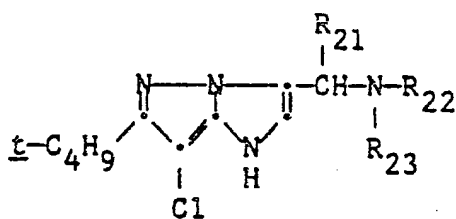


This data demonstrates the shift in hue and increased maximum density provided by the couplers of the invention compared to Comparison A (Compound C-8). The photographic elements of Examples 1 and 2 also provide higher contrast and higher relative speed compared to a photographic element containing comparison coupler A.

Other examples of useful pyrazoloazole couplers are as follows: these can be prepared by methods like

those described above and can be used in a color photographic element as described in Example 1.

Table V



## Example

No.	R <sub>21</sub>	R <sub>22</sub>	R <sub>23</sub>
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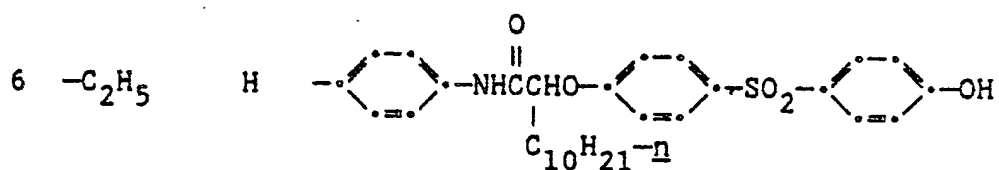
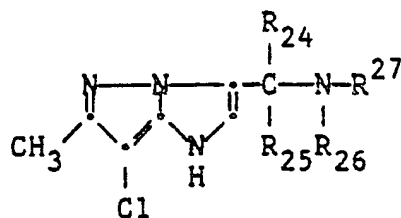


Table V (Continued)

5	7	$-C_2H_5$	$CH_3$	
10	8	$-C_{10}H_{21-n}$	H	
15	9	$-C_{12}H_{25-n}$	H	
20	10	$-C_{12}H_{25-n}$	$C_{12}H_{25-n}$	
25	11	$-C_{12}H_{25-n}$	$CH_3$	
30	12	$-C_{18}H_{37-n}$	H	

Table VI



## Example

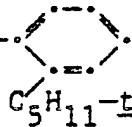





No.	R <sub>24</sub>	R <sub>25</sub>	R <sub>26</sub>	R <sub>27</sub>
13		H	CH <sub>3</sub>	
14			C <sub>2</sub> H <sub>5</sub>	
15		H	C <sub>2</sub> H <sub>5</sub>	
16			C <sub>2</sub> H <sub>5</sub>	C <sub>5</sub> H <sub>10</sub> COOH
17		H		C <sub>18</sub> H <sub>36</sub> COOH
18		H	H	CH <sub>2</sub> CH <sub>2</sub> COOH

Table VI Continued

5	19	$-\text{C}_{12}\text{H}_{25}-\text{n}$	H	$-\text{C}_4\text{H}_9-\text{n}$	$-\text{C}(=\text{O})\text{CH}_2\text{CH}_2\text{COOH}$
10	20	$\text{C}_2\text{H}_5$	$\text{C}_2\text{H}_5$	$\text{C}_{18}\text{H}_{37}-\text{n}$	$-\text{C}(=\text{O})\text{CH}_2\text{COOH}$
15	21	$\text{C}_{10}\text{H}_{21}-\text{n}$	H	H	$-\text{CH}_2-\text{C}_6\text{H}_4-\text{NHSO}_2-\text{C}_6\text{H}_4-\text{COOH}$
20	22	$\text{C}_2\text{H}_5$	H	H	$-\text{CH}_2-\text{C}_6\text{H}_4-\text{NHC}(=\text{O})-\text{C}_6\text{H}_3(\text{O})-\text{CH}(\text{COOH})-\text{C}_{12}\text{H}_{25}-\text{n}$
25	23	$\text{C}_{12}\text{H}_{25}-\text{n}$	$\text{CH}_3$	$\text{C}_{18}\text{H}_{37}-\text{n}$	$-\text{C}(=\text{O})\text{NH}-\text{C}_6\text{H}_4-\text{COOH}$
30	24	$\text{C}_2\text{H}_5$	H	$\text{C}_{18}\text{H}_{37}-\text{n}$	$-\text{C}(=\text{O})\text{NH}-\text{C}_6\text{H}_4-\text{COOH}$
35	25	$\text{C}_2\text{H}_5$	H	$\text{CH}_3$	$-\text{C}(=\text{O})\text{NH}-\text{C}_6\text{H}_4-\text{COOH}$
40	26	$\text{C}_{12}\text{H}_{25}-\text{n}$	H	$\text{C}_4\text{H}_9-\text{i}$	$-\text{C}(=\text{O})\text{CH}_2\text{CH}_2\text{COOH}$
45					
50					
55					



Table VI Continued

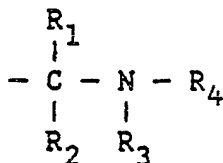
27	$C_2H_5$	H	$\begin{array}{c} -C=O \\   \\ CH_2CH_2COOH \end{array}$	$-(CH_2)_4-O-$		$-C_5H_{11}-t$
28	$C_{12}H_{25}-n$	H	$C_8H_{17}-n$	$\begin{array}{c} -C \\    \\ O \end{array}$	$CH_2CH_2COOH$	
29	$C_2H_5$	H	$C_8H_{17}-n$	$C_8H_{17}-n$		
30	$C_{12}H_{25}-n$	H	$CH_3$			
31	$C_{12}H_{25}-n$	H	$CH_3$		$-OCH_3$	
32	$C_2H_5$	H	$C_{12}H_{25}-n$		$-OH$	
33	$C_{12}H_{25}-n$	H	$C_2H_4OC_2H_4OC_2H_4OCH_3$	$C_2H_4OC_2H_4OC_2H_4OCH_3$		
34	$C_2H_5$	H	$CH_3$		$-CO_2CH_3$	
35	$C_{12}H_{25}-n$	H	$C_2H_4OH$	$C_2H_4OH$		
36	$C_{12}H_{25}-n$	H				
37	$C_{12}H_{25}-n$	H	$CH_3$	$CH_2CH_2CO_2C_2H_5$		

The compounds of the invention of these examples also enable reduced silver interaction.

## Claims

1. A photographic silver halide element comprising a support bearing at least one photographic silver halide emulsion layer and in or adjacent the emulsion layer a pyrazoloazole color coupler having substituted on the azole ring of the coupler a group containing a carbon atom (A) bonded directly to the azole ring and wherein the carbon atom (A) is bonded directly to a nitrogen atom in the group that enables a bathochromic hue shift in a dye formed from the pyrazoloazole coupler.

2. A photographic element as claimed in claim 1 wherein the pyrazoloazole coupler has substituted on the azole ring of the coupler a group represented by the structure:



wherein:

$R_1$  and  $R_2$  are individually hydrogen, unsubstituted or substituted alkyl or unsubstituted or substituted aryl;

$R_3$  and  $R_4$  are individually hydrogen, unsubstituted or substituted alkyl, unsubstituted or

substituted aryl, or  $\begin{array}{c} O \\ || \\ C - R_5 \end{array}$ ; wherein at least one of  $R_3$  and  $R_4$  is alkyl, aryl or

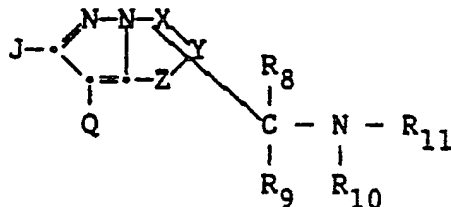
$\begin{array}{c} O \\ || \\ C - R_5 \end{array}$ ;

$R_5$  is unsubstituted or substituted alkyl, unsubstituted or substituted aryl, or  $NH-R_6$ ; and

$R_6$  is unsubstituted or substituted alkyl, unsubstituted or substituted aryl or  $COOR_7$ , wherein  $R_7$  is unsubstituted or substituted alkyl or unsubstituted or substituted aryl.

3. A photographic element as claimed in claim 1 or 2 wherein the pyrazoloazole coupler is a pyrazolotriazole coupler.

4. A photographic element as claimed in claim 1 wherein the pyrazoloazole coupler is represented by the structure:



wherein:

J is a substituent;

Q is hydrogen or a coupling-off group;

X, Y and Z are individually selected from carbon and nitrogen atoms necessary to complete an azole ring;

$R_8$  and  $R_9$  are individually hydrogen, unsubstituted or substituted alkyl or unsubstituted or substituted aryl;

$R_{10}$  and  $R_{11}$  are individually hydrogen, unsubstituted or substituted alkyl;

unsubstituted or substituted aryl or  $\begin{array}{c} O \\ || \\ C - R_{12} \end{array}$ ; wherein at least one of

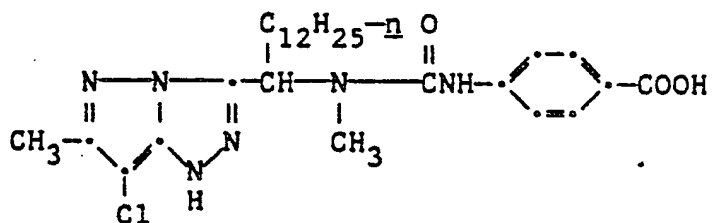
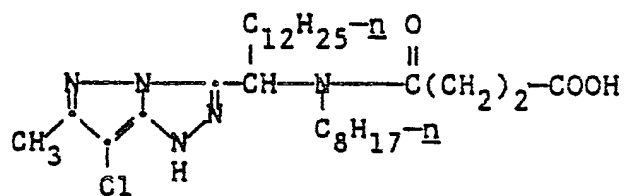
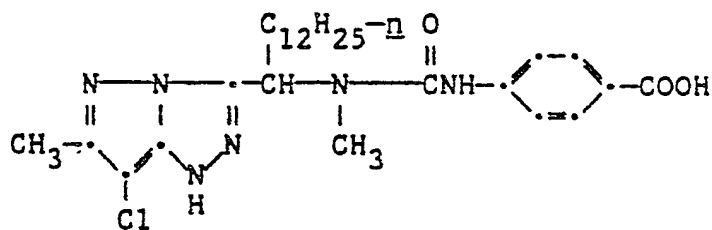
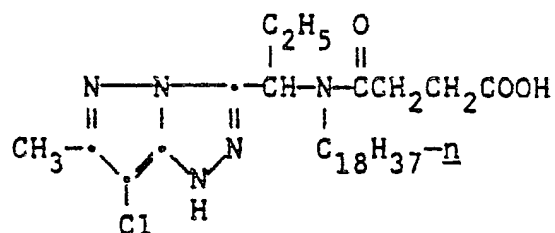
$R_{10}$  and  $R_{11}$  is alkyl, aryl or  $\begin{array}{c} O \\ || \\ C - R_{12} \end{array}$ ;

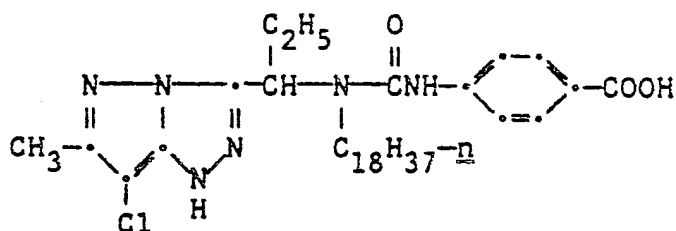
$R_{12}$  is unsubstituted or substituted alkyl, unsubstituted or substituted aryl or  $NH-R_{13}$ ;

$R_{13}$  is unsubstituted or substituted alkyl or unsubstituted or substituted aryl.

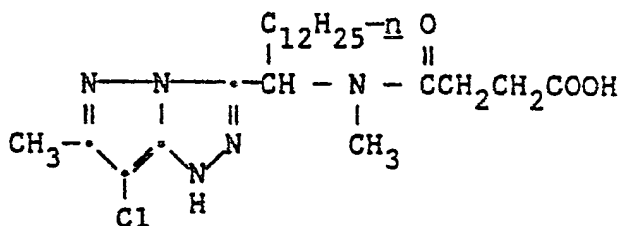
5. A photographic element as claimed in any of claims 1-4 wherein the pyrazoloazole coupler comprises at least one water solubilizing group.

6. A photographic element as claimed in any of claims 1 - 5 wherein the pyrazoloazole coupler is a compound selected from the group consisting of:





and



7. A photographic element as claimed in any of claims 1 - 6 comprising a red-sensitive silver halide emulsion unit having associated therewith a cyan dye image forming material, a green-sensitive silver halide emulsion unit having associated therewith a magenta dye image forming material, and a blue-sensitive silver halide emulsion unit having associated therewith a yellow dye-image forming material, and wherein said pyrazoloazole coupler is associated with at least one of said units.

8. A process of forming a photographic color image which comprises developing an exposed photographic silver halide emulsion layer with a color developing agent in the presence of a pyrazoloazole color coupler as defined in any one of claims 1-6.

9. A photographic magenta dye-forming pyrazoloazole color coupler as defined in any one of claims 1-6.



European Patent  
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# EUROPEAN SEARCH REPORT

Application Number

EP 88 30 2009

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. CL.4)
X	PATENT ABSTRACTS OF JAPAN, vol. 10, no. 232 (P-486)[2288], 12th August 1986; & JP-A-61 65 247 (FUJI PHOTO FILM CO. LTD) 03-04-1986 * Abstract *	1-4	G 03 C 7/38 // C 07 D 487/04
X	EP-A-0 176 804 (FUJI PHOTO FILM) * Claim 1 *	1	
P,X	EP-A-0 226 849 (FUJI PHOTO FILM) * Page 16, compound 22 *	1	
			TECHNICAL FIELDS SEARCHED (Int. CL.4)
			G 03 C C 07 D
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 18-07-1988	Examiner MALHERBE
CATEGORY OF CITED DOCUMENTS			
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document		I : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	

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